

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: December 6, 2002, 16:39:25 : Search time 231.5 Seconds
(without alignment)
14834.978 Million cell updates/sec

Title: US-10-025-514-7
Perfect score: 1525
Sequence: 1 tctagaccatgtcttggaag.....ccaactcagaagttagtcgac 1525

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_101002.*
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24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Query Match	Length	ID	Description
1	1525	100.0	1525	24	ABK88022
2	1196	78.4	1756	24	ABK88023
3	1194.8	78.3	1582	24	ABK88024
4	1191.6	78.1	1525	24	ABK88025
5	1191.4	78.1	1582	24	ABK88027
6	1191.4	78.1	1756	24	ABK88026
7	1182	77.5	1182	24	ABK88015
8	629.4	41.3	1260	19	AAV41730
9	436.4	28.6	1312	16	AAQ89254

10	436.4	28.6	1312	19	AAV28471	Nucleotide sequence
11	436.4	28.6	1312	21	AAZ90199	Human alpha-anti
12	433.4	28.4	1367	22	AA345052	cDNA encoding nove
13	433.2	28.4	1352	13	AAQ31403	Human alpha-1 anti
14	433.2	28.4	1352	24	ABL67511	Thyroid cancer rel
15	433.2	28.4	1371	24	ABR84495	Human cDNA differe
16	433.2	28.4	1371	24	ABL67510	Thyroid cancer rel
17	433.2	28.4	1433	10	AA91077	Sequence encoding
18	433.2	28.4	1434	5	AA40078	Sequence encoding
19	433.2	28.4	1434	20	AA383548	Human alpha-anti-
20	433.2	28.4	5932	21	AA245928	Nucleotide sequenc
21	433.2	28.4	6142	21	AA245932	Nucleotide sequenc
22	433.2	28.4	6142	21	AA245933	Nucleotide sequenc
23	433.2	28.4	6565	21	AA245925	Nucleotide sequenc
24	433.2	28.4	6714	21	AA245930	Nucleotide sequenc
25	433.2	28.4	6924	21	AA245934	Nucleotide sequenc
26	433.2	28.4	6924	21	AA245935	Nucleotide sequenc
27	433.2	28.4	6981	21	AA245931	Nucleotide sequenc
28	433.2	28.4	7054	21	AA245927	Nucleotide sequenc
29	432.8	28.4	7405	21	AA245926	Nucleotide sequenc
30	431.6	28.3	1352	18	AA72858	Nucleotide sequenc
31	430.4	28.2	1185	19	AAV41726	Human alpha-1-anti
32	430	28.2	1434	10	AA90341	Native coding sequ
33	429.6	28.2	1312	10	AA97127	Sequence of alpha-
34	429	28.1	1189	13	AAQ21125	Alpha-1-antitrypsi
35	428.4	28.1	1378	13	AAQ23746	Alpha-1-antitrypsi
36	428.4	28.1	1396	11	AAQ03184	Entire sequence of
37	426.8	28.0	1185	7	AA60417	Human alpha 1-anti
38	426.8	28.0	1423	6	AA50540	Sequence encoding
39	425.2	27.9	1299	6	AA50540	Sequence of human
40	425.2	27.9	1378	6	AA50021	Sequence encoding
41	411.2	27.0	1390	22	AAH23089	Osteoarthritis tis
42	401.4	26.3	2013	24	ABL59152	Sequence of fusion
43	375.6	24.6	1242	18	AA79493	Protease inhibitor
44	362.8	23.8	1242	18	AA78180	Recombinant squirr
45	360.8	23.7	1312	10	AA91078	Alpha-1-antitrypsi

ALIGNMENTS

RESULT 1
ABK88022
ID ABK88022 standard; DNA; 1525 BP.
XX
AC ABK88022;
XX
DT 07-OCT-2002 (first entry)
XX
DE DNA sequence encoding SLAP1 fusion protein.
XX
KW Yeast; alpha factor; gene; ds; Alzheimer's disease; SLAP1;
KW malaria; emphysema; asthma; chronic obstructive pulmonary disease;
KW cystic fibrosis; otitis media; otitis externa; HIV; psoriasis; eczema;
KW human immunodeficiency virus; atopic dermatitis; muscular dystrophy;
KW herpes; ulceration; sepsis; rheumatoid arthritis; periodontal disease;
KW tumour metastasis; tumour angiogenesis; osteoporosis; Paget's disease;
KW glomerulonephritis; scleroderma; hypertension.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT RBS 6..8
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FT /standard_name= "Ribosome binding site"
FT CDS 9..1520
FT /tag= b
FT /product= "SLAP1 fusion protein"
FT misc_feature 12..332
FT /tag= c
FT /note= "SLP1 coding region"
FT misc_feature 333-335

FT misc_feature /tag= d /note= "linking codon"
FT 336..1517 /tag= e /note= "AAT coding region"
FT 336..1517 /tag= e /note= "AAT coding region"

W0200250287-A2.

27-JUN-2002.

18-DEC-2001; 2001WO-US49256.

18-DEC-2000; 2000US-256699P.

20-NOV-2001; 2001US-331966P.

(ARRI-) ARRIVA PHARM INC.

Barr PJ, Gibson HL, Pemberton P;

WPI; 2002-500631/53.

P-PSDB; RAU99881.

Novel fusion protein useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, comprises a first protease inhibitor comprising alpha 1-antitrypsin and a second protease inhibitor -

Example 1; Page 73-73; 134pp; English.

This invention relates to a novel fusion protein comprising a first protease inhibitor comprising an alpha 1-antitrypsin or its functionally active portion and a second protease inhibitor or its functionally active portion. The fusion proteins of the invention may act as an inhibitor of protease activity. The fusion protein of the invention is useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, chronic obstructive pulmonary disease, cystic fibrosis, otitis media, otitis externa or HIV infection, or for treating an individual suffering from or at risk for a disease or disorder involving unwanted protease activity. The proteins are useful for treating dermatological diseases such as atopic dermatitis, eczema and psoriasis, in inflammatory responses to viral infection, and for treating herpes infection, corneal or epidermal ulceration, chronic non-healing wounds, sepsis, rheumatoid arthritis, periodontal disease, tumour metastasis and tumour angiogenesis, gastric ulceration, osteoporosis, Paget's disease, glomerulonephritis, scleroderma, bacterial infection, Alzheimer's disease, hypertension and muscular dystrophy. The present sequence represents the DNA encoding the SLAP1 fusion protein of the invention.

Sequence 1525 BP; 467 A; 286 C; 314 G; 458 T; 0 other;

Query Match 100.0%; Score 1525; DB 24; Length 1525;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1525; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 1 TCTAGACCATGCTGGAAGTCTTTCAAGCCCGGTGTTGTCACCAAGAGAGTCCGCTC 60
QY 61 AATGTTTGAGATACAGAAGCCAGAATGTCAATCCGACTGGCAATGTCACAGGTAAGA 120
DB 61 AATGTTTGAGATACAGAAGCCAGAATGTCAATCCGACTGGCAATGTCACAGGTAAGA 120
QY 121 GATGTTGCCAGACACTTGGGTATCAAGTGTCTAGACCCAGTGTGACACCCCAACCCCA 180
DB 121 GATGTTGCCAGACACTTGGGTATCAAGTGTCTAGACCCAGTGTGACACCCCAACCCCA 180
QY 181 CTAGAAGAACCCAGGTAAGTGTCCAGTCTACTTACGGTCAATGTTGATGTTGAACCCAC 240
DB 181 CTAGAAGAACCCAGGTAAGTGTCCAGTCTACTTACGGTCAATGTTGATGTTGAACCCAC 240
QY 241 CAAACTTCTGTGAATGACCGGTCAATGTAAGAGAGACTTGAAGTGTGATGGGTAGT 300
DB 241 CAAACTTCTGTGAATGACCGGTCAATGTAAGAGAGACTTGAAGTGTGATGGGTAGT 300

DB 241 CAAACTTCTGTGAATGACCGGTCAATGTAAGAGAGACTTGAAGTGTGATGGGTAGT 300
QY 301 GTGGTAAGTCTCTGTGTTTCCCGAGTCAAGGCGCATGGAAGACCTCAAGGCGACGCGCTC 360
DB 301 GTGGTAAGTCTCTGTGTTTCCCGAGTCAAGGCGCATGGAAGACCTCAAGGCGACGCGCTC 360
QY 361 AAAAAACGACACCAAGTCAATCAGACCAAGCAATCCGACTTTTAAATAAAATTAATCTCAA 420
DB 361 AAAAAACGACACCAAGTCAATCAGACCAAGCAATCCGACTTTTAAATAAAATTAATCTCAA 420
QY 421 AATTAGCCGAATTTGCTTTTCTTTGATAGACAATTAGCTCATCAAAAGTAATTTCTACTA 480
DB 421 AATTAGCCGAATTTGCTTTTCTTTGATAGACAATTAGCTCATCAAAAGTAATTTCTACTA 480
QY 481 ACATTTTCTAGTCTGTTTCTTATGCGCACTGCTTTTCCGCCATGTTGAGTTAGTACTA 540
DB 481 ACATTTTCTAGTCTGTTTCTTATGCGCACTGCTTTTCCGCCATGTTGAGTTAGTACTA 540
QY 541 AAGCCGATACCCATGACGAGATTTTGAAGGTTTAAACTTTTAAATTTGACCGCAATCCAG 600
DB 541 AAGCCGATACCCATGACGAGATTTTGAAGGTTTAAACTTTTAAATTTGACCGCAATCCAG 600
QY 601 AAGCCCAATTCAGAGGTTTTCAGAGTGTGTTGAGAATTTGAATCAACCTGATTTCTC 660
DB 601 AAGCCCAATTCAGAGGTTTTCAGAGTGTGTTGAGAATTTGAATCAACCTGATTTCTC 660
QY 661 AATTGCAATTAACCTACTGTTAAGCGTTTATTTTGTCTGAAGGTTTAAATTTGAGTGA 720
DB 661 AATTGCAATTAACCTACTGTTAAGCGTTTATTTTGTCTGAAGGTTTAAATTTGAGTGA 720
QY 721 AATTCTTAGAGAGCGTCAAGAACTATATCATAGTAGGCTTTTACCGTTAAATTTGGTG 780
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QY 781 ATACTGAGGAGCTTAAAGCAAAATTAATGATTTGTTGAGAAGGCGACCGGTAAGA 840
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QY 901 TTTTCAAGGTTAAGTGGGAGCGTCTTTCGAGGTTAAAGATACCTGAAGAGAGAGATTTTC 960
DB 901 TTTTCAAGGTTAAGTGGGAGCGTCTTTCGAGGTTAAAGATACCTGAAGAGAGAGATTTTC 960
QY 961 ATGTTGATCAAGTTACTACTGTCAAAGTTCCATGATGAAAAGAGCTGGGTATGTTCAATA 1020
DB 961 ATGTTGATCAAGTTACTACTGTCAAAGTTCCATGATGAAAAGAGCTGGGTATGTTCAATA 1020
QY 1021 TTCAACATTCGCAAAATAAATTAAGTCTTTGGGTCTTTATTAATGAAGTATTTAGGTAA 1080
DB 1021 TTCAACATTCGCAAAATAAATTAAGTCTTTGGGTCTTTATTAATGAAGTATTTAGGTAA 1080
QY 1081 CTGCTATTTTCTTACCGAGCAAGTAAAGTCTTCAACATTTAGAGAAATGAGTTGACTC 1140
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QY 1141 ATGACATATTTACTAAATTTTATAGAGAACGAGGATCGTGGTACCGCTTCTCTGCACCTGC 1200
DB 1141 ATGACATATTTACTAAATTTTATAGAGAACGAGGATCGTGGTACCGCTTCTCTGCACCTGC 1200
QY 1201 CAAAGTTAAGTATACCGGTACTTACGACTTAAATCTGTTTAGGCCAGTTAGGTATTA 1260
DB 1201 CAAAGTTAAGTATACCGGTACTTACGACTTAAATCTGTTTAGGCCAGTTAGGTATTA 1260
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DB 1261 CCAAGTCTTTTCTAACGTCGCGATTTGAGTGGTGTACTGAAGAAGCTCCATTTAAAT 1320
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1221 CCAATGATGAAAGACTGGGTATGTTCAATATTCACATTCGAAAATTAAGTCTTGG 1280
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Db 1281 GTCTTATTAATGAAGTATTAGGTAAAGCTACTGCTATTTTTTTTCCAGACGAAGGT 1340
QY 1110 AAGCTTCAACATTTAGAGAATAGTGTGACTCATGACATTAATTAATAATTTTACAGAAC 1169
Db 1341 AAGCTTCAACATTTAGAGAATAGTGTGACTCATGACATTAATTAATAATTTTACAGAAC 1400
QY 1170 GAGATCGTGGTACGCTCTCTGACCTGCCAAAGTTAAGTATACACCGGTACTTACGAC 1229
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QY 1230 TAAATCTGTTTATAGCCAGTTAGTATTACCAAGTTTTTTTCTAACGGTCCGATTTG 1289
Db 1461 TTAATCTGTTTATAGCCAGTTAGTATTACCAAGTTTTTTTCTAACGGTCCGATTTG 1520
QY 1290 AGTGTGTTTACTGAAGAAGCTCCATTAATTTAGTAAAGCTGTTCACAAAGCCGCTTTA 1349
Db 1521 AGTGTGTTTACTGAAGAAGCTCCATTAATTTAGTAAAGCTGTTCACAAAGCCGCTTTA 1580
QY 1350 ACTATTGATGAAAGGTTACCGAGCGCGCGCTATGTTCTGGAGCTATTCCTCAATG 1409
Db 1581 ACTATTGATGAAAGGTTACCGAGCGCGCGCTATGTTCTGGAGCTATTCCTCAATG 1640
QY 1410 AGCATTCACAGAAAGTTAAATTTAATAACCATTCGTTTTCTGATGATCGAGCAGAAC 1469
Db 1641 AGCATTCACAGAAAGTTAAATTTAATAACCATTCGTTTTCTGATGATCGAGCAGAAC 1700
QY 1470 ACTAAAGCCCATGTTTATGGTAAAGTTGTCAACCAACTCAGAAAGTAGTCGAC 1525
Db 1701 ACTAAAGCCCATGTTTATGGTAAAGTTGTCAACCAACTCAGAAAGTAGTCGAC 1756

RESULT 3
ABK88024
ID ABK88024 standard; DNA; 1582 BP.
XX AC ABK88024;
XX DT 07-OCT-2002 (first entry)
XX DE DNA sequence encoding N-TAP1 fusion protein.
XX KW NTAP1; gene; ds; Alzheimer's disease; tumour angiogenesis;
XX KW malaria; emphysema; asthma; chronic obstructive pulmonary disease;
XX KW cystic fibrosis; otitis media; otitis externa; HIV; psoriasis; eczema;
XX KW human immunodeficiency virus; atopic dermatitis; muscular dystrophy;
XX KW herpes; ulceration; sepsis; rheumatoid arthritis; periodontal disease;
XX KW tumour metastasis; osteoporosis; Paget's disease; scleroderma;
XX KW glomerulonephritis; hypertension.
XX OS Homo sapiens.
XX OS Synthetic.
XX PH Key
XX FT RBS Location/Qualifiers
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FT 390..392 /*tag= d
FT /*note= "linking codon"
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FT /*note= "AAT coding region"
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XX WO200250287-A2.
XX PD 27-JUN-2002.
XX PF 18-DEC-2001; 2001WO-US49256.
XX PR 18-DEC-2000; 2000US-256699P.
XX PR 20-NOV-2001; 2001US-331966P.
XX PA (ARRI-) ARRIVA PHARM INC.
XX PI Barr PJ, Gibson HL, Pemberton P;
XX DR WPI; 2002-500631/53.
XX DR P-PSDB; AAU99883.
XX Novel fusion protein useful for inhibiting protease activity associated
XX with a disorder such as emphysema, asthma, comprises a first protease
XX inhibitor comprising alpha 1-antitrypsin and a second protease
XX inhibitor -
XX Example 2; Page 85-86; 134pp; English.
XX This invention relates to a novel fusion protein comprising a first
XX protease inhibitor comprising an alpha 1-antitrypsin or its functionally
XX active portion and a second protease inhibitor or its functionally
XX active portion. The fusion proteins of the invention may act as an
XX inhibitor of protease activity. The fusion protein of the invention
XX is useful for inhibiting protease activity associated with a disorder
XX such as emphysema, asthma, chronic obstructive pulmonary disease,
XX cystic fibrosis, otitis media, otitis externa or HIV infection, or
XX for treating an individual suffering from or at risk for a disease or
XX disorder involving unwanted protease activity. The proteins are useful
XX for treating dermatological diseases such as atopic dermatitis, eczema
XX and psoriasis, in inflammatory responses to viral infection, and for
XX treating herpes infection, corneal or epidermal ulceration, chronic
XX non-healing wounds, sepsis, rheumatoid arthritis, periodontal disease,
XX tumour metastasis and tumour angiogenesis, gastric ulceration,
XX osteoporosis, Paget's disease, glomerulonephritis, scleroderma, malaria,
XX bacterial infection, Alzheimer's disease, hypertension and muscular
XX dystrophy. The present sequence represents the DNA encoding the
XX NTAP1 fusion protein of the invention.
XX SQ Sequence 1582 BP; 464 A; 333 C; 329 G; 456 T; 0 other;
XX Query Match 78.3%; Score 1194.8; DB 24; Length 1582;
XX Best Local Similarity 99.8%; Pred. No. 2.7e-290;
XX Matches 1196; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 328 AGGCAATGGAAGACCCCTCAAGGCGCGCGCTCAAAAACCGACACCATCATCAGACC 387
Db 385 AGGAAATGGAAGACCCCTCAAGGCGCGCGCTCAAAAACCGACACCATCATCAGACC 444
QY 388 AAGACCATCCGACTTTTAAATAAATTAATTAATTAATTAATTAATTAATTAATTAAT 447
Db 445 AAGACCATCCGACTTTTAAATAAATTAATTAATTAATTAATTAATTAATTAATTAAT 504
QY 448 ATAGACAATTAAGTCATCAAGTAATTCATCAATTTTCTAGTCTCTGTTCTATTG 507
Db 505 ATAGACAATTAAGTCATCAAGTAATTCATCAATTTTCTAGTCTCTGTTCTATTG 564
QY 508 CCACTGCTTCGCCATGTTGAGTTAGTACTAAAGCCGATACCCATGACGAGATTTAG 567
Db 565 CCACTGCTTCGCCATGTTGAGTTAGTACTAAAGCCGATACCCATGACGAGATTTAG 624
QY 568 AAGGTTTAAACTTTAATTTGACCGAAATCCAGAACCCCAATTCAGGAGGGTTTCAAG 627
Db 625 AAGGTTTAAACTTTAATTTGACCGAAATCCAGAACCCCAATTCAGGAGGGTTTCAAG 684
QY 628 AGTTGTTGAGAACTTTGAATCAACTGATTTCTCAATTTGCAATTAACCTACTGTTAAGCGTT 687
Db 685 AGTTGTTGAGAACTTTGAATCAACTGATTTCTCAATTTGCAATTAACCTACTGTTAAGCGTT 744

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QY 688 TATTTTCTCTGAAGGTTTAAATTTGGTTGACAAATTCCTAGAGACGCTCAAGAACTAT 747
KW |
XX |
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Db 745 TATTTTCTCTGAAGGTTTAAATTTGGTTGACAAATTCCTAGAGACGCTCAAGAACTAT 804
QY 748 ATCATAGTGAAGGCTTTTACCGTTAAATTTGGTGATCTAGGAGAGCTAAAGCAAAATTA 807
DQ |
Db 805 ATCATAGTGAAGGCTTTTACCGTTAAATTTGGTGATCTAGGAGAGCTAAAGCAAAATTA 864
QY 808 ATGATTATGTTGAAGAGCCAGGCTAAGATCGTTGACCTAGTTAAAGAAATTTAGATC 867
DQ |
Db 865 ATGATTATGTTGAAGAGCCAGGCTAAGATCGTTGACCTAGTTAAAGAAATTTAGATC 924
QY 868 GTGATACCGCTCTTCGACACTAGTTAACTATATTTTTCAGGGTAAGTGGGAAGCTCCTT 927
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QY 1108 GTAGCTTCAACATTTAGAGATGAGTGTGATCATGATGATGATGATGATGATGATGAT 1167
DQ |
Db 1165 GTAGCTTCAACATTTAGAGATGAGTGTGATCATGATGATGATGATGATGATGATGAT 1224
QY 1168 ACAGGATCGTGTGAGGCTTCTCTGACCTGCGCAAGTTAAGTATCACCGGTACTTACG 1227
DQ |
Db 1225 ACAGGATCGTGTGAGGCTTCTCTGACCTGCGCAAGTTAAGTATCACCGGTACTTACG 1284
QY 1228 ACTTAAATCTGTTTAGGCGAGTTAGTATTAACCAAGTTTTCCTTACCGGTGCGGAT 1287
DQ |
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QY 1348 TAATCTATGATGAAGAGGTACGAGCGCGCGCGCTATGTTCTCGGAAGCTATTCCAA 1407
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QY 1408 TGAGCATTTCCACCAAGAGTTAATTTAATTAACCATTCGTTTTCGTGATGATGATGATGAT 1467
DQ |
Db 1465 TGAGCATTTCCACCAAGAGTTAATTTAATTAACCATTCGTTTTCGTGATGATGATGATGAT 1524
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DQ |
Db 1525 ACATAAAGCCCATTTGTTTATGGTAAAGTGTGTCACCAACCTCAGAGTGTAGTCGAC 1582
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RESULT 4

ABK88025

ID ABK88025 standard; DNA: 1525 BP.

XX ABK88025;

AC ABK88025;

DT 07-OCT-2002 (first entry)

DE DNA sequence encoding rSLAP1 fusion protein.

XX rSLAP1; gene: ds; Alzheimer's disease; tumour angiogenesis;
KW malaria; emphysema; asthma; chronic obstructive pulmonary disease;
KW cystic fibrosis; otitis media; otitis externa; HIV; psoriasis; eczema;
KW human immunodeficiency virus; atopic dermatitis; muscular dystrophy;
KW herpes; ulceration; sepsis; rheumatoid arthritis; periodontal disease;

KW tumour metastasis; osteoporosis; Paget's disease; scleroderma;
KW glomerulonephritis; hypertension.

OS Homo sapiens.

XX Synthetic.

FH Key

FT RBS

FT Location/Qualifiers

FT 6..8

FT /tag= a

FT /standard_name= "Ribosome binding site"

FT 9..1520

FT /tag= b

FT /product= "rSLAP1 fusion protein"

FT 12..1193

FT /tag= c

FT /note= "AAT coding region"

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FT /note= "Linking codon"

FT 1197..1517

FT /tag= e

FT /note= "SLP1 coding region"

FT WO200250287-A2.

XX 27-JUN-2002.

XX 18-DEC-2001; 2001WO-US49256.

XX 18-DEC-2000; 2000US-256699P.

XX 20-NOV-2001; 2001US-331966P.

XX (ARRI-) ARRIVA PHARM INC.

XX Barr PJ, Gibson HL, Pemberton P;

XX WPI: 2002-500631/53.

XX P-PSDB; AAU99884.

Novel fusion protein useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, comprises a first protease inhibitor comprising alpha 1-antitrypsin and a second protease inhibitor.

Example 3; Page 89-90; 134pp; English.

This invention relates to a novel fusion protein comprising a first protease inhibitor comprising an alpha 1-antitrypsin or its functionally active portion and a second protease inhibitor or its functionally active portion. The fusion proteins of the invention may act as an inhibitor of protease activity. The fusion protein of the invention is useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, chronic obstructive pulmonary disease, cystic fibrosis, otitis media, otitis externa or HIV infection, or for treating an individual suffering from or at risk for a disease or disorder involving unwanted protease activity. The proteins are useful for treating dermatological diseases such as atopic dermatitis, eczema and psoriasis, in inflammatory responses to viral infection, and for treating herpes infection, corneal or epidermal ulceration, chronic non-healing wounds, sepsis, rheumatoid arthritis, periodontal disease, tumour metastasis and tumour angiogenesis, gastric ulceration, osteoporosis, Paget's disease, glomerulonephritis, scleroderma, malaria, bacterial infection, Alzheimer's disease, hypertension and muscular dystrophy. The present sequence represents the DNA encoding the rSLAP1 fusion protein of the invention.

Sequence 1525 BP; 467 A; 287 C; 314 G; 457 T; 0 other;

Query Match

Best Local Similarity 78.1%; Score 1191.6; DB 24; Length 1525;

Matches 1194; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 325 TCAAGGCCATGGGAAGACCTCAAGGCGCGCTCAAAAAACCCAGCAGTCATCAGC 384

1081 CAATGAGCATCCACCAGAAAGTTAAATTAATAAACCAATTCGTTTTCTGTATGATCGAGC 1140
 1465 AGAACACTAAAGCCCATTTGTTATGGGTAAAGTTGTCAACCCAACTCAGAAGTAGTC 1522
 1141 AGAACACTAAAGCCCATTTGTTATGGGTAAAGTTGTCAACCCAACTCAGAAGTAGTC 1198

RESULT 5
 ABK88027
 ID ABK88027 standard; DNA; 1582 BP.
 XX
 AC ABK88027;
 XX
 XX 07-OCT-2002 (first entry)
 XX
 XX DNA sequence encoding rN-TAP1 fusion protein.

rN-TAP1; gene; ds; Alzheimer's disease; tumour angiogenesis;
 malaria; emphysema; asthma; chronic obstructive pulmonary disease;
 cystic fibrosis; otitis media; otitis externa; HIV; psoriasis; eczema;
 human immunodeficiency virus; atopic dermatitis; muscular dystrophy;
 herpes; ulceration; sepsis; rheumatoid arthritis; periodontal disease;
 tumour metastasis; osteoporosis; Paget's disease; scleroderma;
 glomerulonephritis; hypertension.

Homo sapiens.
 OS Synthetic.

XX Key Location/Qualifiers
 XX RBS 6..8
 FT /tag= a
 FT /standard_name= "Ribosome binding site"
 FT CDS 9..1577
 FT /tag= b
 FT /product= "rTAP1 fusion protein"
 FT 12..1193
 FT /tag= c
 FT /note= "AAT coding region"
 FT 1194..1196
 FT /tag= d
 FT /note= "linking codon"
 FT 1197..1574
 FT /tag= e
 FT /note= "TIMP-1 coding region"

WO200250287-A2.

27-JUN-2002.

18-DEC-2001; 2001WO-US49256.

18-DEC-2000; 2000US-256699P.

20-NOV-2001; 2001US-331966P.

(ARRI-) ARRIVA PHARM INC.

Barr PJ, Gibson HL, Pemberton P;

WPI; 2002-500631/53.

P-PSDB; AAU99885.

Novel fusion protein useful for inhibiting protease activity associated
 with a disorder such as emphysema, asthma, comprises a first protease
 inhibitor comprising alpha 1-antitrypsin and a second protease
 inhibitor

Example 3; Page 95-96; 134pp; English.

This invention relates to a novel fusion protein comprising a first
 protease inhibitor comprising an alpha 1-antitrypsin or its functionally
 active portion and a second protease inhibitor or its functionally
 active portion. The fusion proteins of the invention may act as an
 inhibitor of protease activity. The fusion protein of the invention

1 TCTAGACCATGGAAGACCTCCTAAGCGGAGCGGCTCAAAAAACGACACCATGATCATCAG 60
 385 ACCAAGACCATCGGACTTTTATAAAATTAACCTCAAAATTTAGCCGAATTTGCTTTTCTT 444
 61 ACCAAGACCATCGGACTTTTATAAAATTAACCTCAAAATTTAGCCGAATTTGCTTTTCTT 120
 445 TGTATAGACAATTAGCTCATCAAGTAATTTCTACTAACAATTTTGTAGTCTGTTTCTTA 504
 121 TGTATAGACAATTAGCTCATCAAGTAATTTCTACTAACAATTTTGTAGTCTGTTTCTTA 180
 505 TTGCCACTGCTTTCGCCATGTTGAGTTTACTTAAGCCGATACCCATGACGAGATTT 564
 181 TTGCCACTGCTTTCGCCATGTTGAGTTTACTTAAGCCGATACCCATGACGAGATTT 240
 565 TAGAAGGTTTAAACTTTAAATTTGACCGAATCCGAGGCCCAAAATTCACGAGGTTTTC 624
 241 TAGAAGGTTTAAACTTTAAATTTGACCGAATCCGAGGCCCAAAATTCACGAGGTTTTC 300
 625 AAGAGTTGTTGAGAACCTTTGAATCAACCTGATTTCTCAATTTGCAATTAACCTACTGCTAAG 684
 301 AAGAGTTGTTGAGAACCTTTGAATCAACCTGATTTCTCAATTTGCAATTAACCTACTGCTAAG 360
 685 GTTATATTTTCTGCTGAAGGTTTAAATTTGCTGACAAATTTCTAGAACGCTCAAGAAC 744
 361 GTTATATTTTCTGCTGAAGGTTTAAATTTGCTGACAAATTTCTAGAACGCTCAAGAAC 420
 745 TATATCATAGTAGGCTTTTACCGTTAAATTTTGGTGATCTAGGAGGCTAAAGACAA 804
 421 TATATCATAGTAGGCTTTTACCGTTAAATTTTGGTGATCTAGGAGGCTAAAGACAA 480
 805 TTAATGATTATGTTGAGAACGCTTAAAGGCTGAGTAAAGTAAAGTAAAGTAAAGTAAAG 864
 481 TTAATGATTATGTTGAGAACGCTTAAAGGCTGAGTAAAGTAAAGTAAAGTAAAGTAAAG 540
 865 ATCGTGATACCGTCTTCCGACTAGTTAACTATATTTTTCAGGGTAAAGTAAAGTAAAGT 924
 541 ATCGTGATACCGTCTTCCGACTAGTTAACTATATTTTTCAGGGTAAAGTAAAGTAAAGT 600
 925 CTTTCGAGGTTAAAGATTAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAG 984
 601 CTTTCGAGGTTAAAGATTAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAG 660
 985 AAGTTCCCAATGATGAAAGCTGGGTATGTTCAATATTCACATTCGAAATTAAGTT 1044
 661 AAGTTCCCAATGATGAAAGCTGGGTATGTTCAATATTCACATTCGAAATTAAGTT 720
 1045 CTTGGGCTCTTAATTAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAG 1104
 721 CTTGGGCTCTTAATTAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAGTAAAG 780
 1105 AAGTTAAAGCTTCAACATTTAGAGATGAGTTGACTCATGACATTTATTAATTAATTTTAC 1164
 781 AAGTTAAAGCTTCAACATTTAGAGATGAGTTGACTCATGACATTTATTAATTAATTTTAC 840
 1165 AGAACGAGGATCGTCTGAGCGGCTCTCTGACCTGCCAAAGTTAAGTATCACCGGACTTT 1224
 841 AGAACGAGGATCGTCTGAGCGGCTCTCTGACCTGCCAAAGTTAAGTATCACCGGACTTT 900
 1225 ACAGCTTAAATCTGTTTTCAGCCAGTTAGGTATTTACCAAGTTTTCCTCAACGCTGCCG 1284
 901 ACAGCTTAAATCTGTTTTCAGCCAGTTAGGTATTTACCAAGTTTTCCTCAACGCTGCCG 960
 1285 ATTTGAGTGTGTTTACTGAAGAGCTCCATTAAATTTGAGTAAAGTGTTCACAAAGCGG 1344
 961 ATTTGAGTGTGTTTACTGAAGAGCTCCATTAAATTTGAGTAAAGTGTTCACAAAGCGG 1020
 1345 TCTTAATATGATGAAAGGTTACCGAGGCGCGCGCTATGTTCTCTGAAAGCTATTC 1404
 1021 TCTTAATATGATGAAAGGTTACCGAGGCGCGCGCTATGTTCTCTGAAAGCTATTC 1080
 1405 CAATGAGCATTCACACAGAGTTAAATTTAATAAACCAATTCGTTTTCTGTATGATCGAGC 1464

21 CTTGGGCTTATTAAATGAAGTATTTAGGTAAAGCTACTGCTATTATTTTATTTTACGACG 780

...GGGTTTAAAGCGCTACTCGTATTTTTTTTACCAGACG 780

WO200250287-A2.

27-JUN-2002.
 18-DEC-2001; 2001WO-US49256.
 18-DEC-2000; 2000US-256699P.
 20-NOV-2001; 2001US-331966P.
 (ARRI-) ARRIVA PHARM INC.
 Barr PJ, Gibson HL, Pemberton P;
 WPI; 2002-500631/53.
 P-PSDB; AAU99889.
 Novel fusion protein useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, comprises a first protease inhibitor comprising alpha 1-antitrypsin and a second protease inhibitor -
 Example 3; Page 92-93; 134pp; English.
 This invention relates to a novel fusion protein comprising a first protease inhibitor comprising an alpha-1-antitrypsin or its functionally active portion and a second protease inhibitor or its functionally active portion. The fusion proteins of the invention may act as an inhibitor of protease activity. The fusion protein of the invention is useful for inhibiting protease activity associated with a disorder such as emphysema, asthma, chronic obstructive pulmonary disease, cystic fibrosis, otitis media, otitis externa or HIV infection, or for treating an individual suffering from or at risk for a disease or disorder involving unwanted protease activity. The proteins are useful for treating dermatological diseases such as atopic dermatitis, eczema and psoriasis, in inflammatory responses to viral infection, chronic treating herpes infection, corneal or epidermal ulceration, chronic non-healing wounds, sepsis, rheumatoid arthritis, periodontal disease, tumour metastasis and tumour angiogenesis, gastric ulceration, osteoporosis, Paget's disease, glomerulonephritis, scleroderma, malaria, bacterial infection, Alzheimer's disease, hypertension and muscular dystrophy. The present sequence represents the DNA encoding the rTAP1 fusion protein of the invention.
 Sequence 1756 BP; 493 A; 394 C; 374 G; 495 T; 0 other;
 Query Match 78.1%; Score 1191.4; DB 24; Length 1756;
 Best Local Similarity 99.5%; Pred. No. 2e-289;
 Matches 1195; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
 325 TCAAGGCCATGGAGACCCCTCAAGCGACGCGCTCAAAAACCGACACCGATCATCAGC 384
 1 TCTAGACCATGGAGACCCCTCAAGCGACGCGCTCAAAAACCGACACCGATCATCAGC 60
 365 ACCAAGACCATCCGACTTTTAAATAAAATTACTCCAAATTTAGCCGAATTTGCTTTTCTT 444
 61 ACCAAGACCATCCGACTTTTAAATAAAATTACTCCAAATTTAGCCGAATTTGCTTTTCTT 120
 445 TGTATAGACAATAGCTCATCAAGTAATCTACTACATTTTCTTCTGCTTTCTTCTA 504
 121 TGTATAGACAATAGCTCATCAAGTAATCTACTACATTTTCTTCTGCTTTCTTCTA 180
 505 TTGCCACTGCTTTGCCCATGTTGAGTTTAGTACTAAAGCCGATACCATGACGAGATTT 564
 181 TTGCCACTGCTTTGCCCATGTTGAGTTTAGTACTAAAGCCGATACCATGACGAGATTT 240
 565 TAGAAGGTTTAAACTTTTAAATTTAGCCGAATTTCCAGAACCCCAATTCACGAGGTTTC 624
 241 TAGAAGGTTTAAACTTTTAAATTTAGCCGAATTTCCAGAACCCCAATTCACGAGGTTTC 300
 625 AGAGTTGTTGAGAACTTTGATCAACCTGATTTCTCAATTCGAATTAACCTAGTGAACG 684
 301 AGAGTTGTTGAGAACTTTGATCAACCTGATTTCTCAATTCGAATTAACCTAGTGAACG 360
 685 GTTATTTTGTCTGAAGGTTTAAATTTGAGTTTAAATTTGAGTTTAAATTTGAGTTTAAATTT 744

Db 361 GTTATTTTGTCTGAAGGTTTAAATTTGGTTACAAATTCCTAGAGACGCTCAAGAAC 420
 QY 745 TATATCATAGTGAAGGCTTTTACCGTTAAATTTGGTGATACTGAGGAGCTAAAGACAAA 804
 Db 421 TATATCATAGTGAAGGCTTTTACCGTTAAATTTGGTGATACTGAGGAGCTAAAGACAAA 480
 QY 805 TTAATGATTTATGTTGAGAAAGGACCCAGGTAAGATCGTTGACCTAGTAAAGAAATTAG 864
 Db 481 TTAATGATTTATGTTGAGAAAGGACCCAGGTAAGATCGTTGACCTAGTAAAGAAATTAG 540
 QY 865 ATCGTGATACCGCTTTCGCACTAGTTAACTATATATTTTTCAGGGTAAGTGGGAACGTC 924
 Db 541 ATCGTGATACCGCTTTCGCACTAGTTAACTATATATTTTTCAGGGTAAGTGGGAACGTC 600
 QY 925 CTTTCGAGGTTTAAAGTACTGAGAGAGAAATTTTCAATGTTGATCAAGTTTACTACTGCTCA 984
 Db 601 CTTTCGAGGTTTAAAGTACTGAGAGAGAAATTTTCAATGTTGATCAAGTTTACTACTGCTCA 660
 QY 985 AAGTTCGAATGATGAAAGACTGGGTATGTTCAATATCAACATTCGCAAAATTAAGTT 1044
 Db 661 AAGTTCGAATGATGAAAGACTGGGTATGTTCAATATCAACATTCGCAAAATTAAGTT 720
 QY 1045 CTTTCGAGGTTTAAAGTACTGAGAGAGAAATTTTCAATGTTGATCAAGTTTACTACTGCTCA 1104
 Db 721 CTTTCGAGGTTTAAAGTACTGAGAGAGAAATTTTCAATGTTGATCAAGTTTACTACTGCTCA 780
 QY 1105 AAGTAAAGCTTCAACATTTAGAGATGAGTTGACTCATGACATATTACTAAATTTTACTAG 1164
 Db 781 AAGTAAAGCTTCAACATTTAGAGATGAGTTGACTCATGACATATTACTAAATTTTACTAG 840
 QY 1165 AGAACGAGATCGTGTAGCGCTTCTCTGCACCTGCCAAAGTTAAGTATCACCGGTACTT 1224
 Db 841 AGAACGAGATCGTGTAGCGCTTCTCTGCACCTGCCAAAGTTAAGTATCACCGGTACTT 900
 QY 1225 ACAGCTTAAATCTGTTTGTAGCCAGTATGAGTATTAACAAAGTTTCTTCAACGGTCCG 1284
 Db 901 ACAGCTTAAATCTGTTTGTAGCCAGTATGAGTATTAACAAAGTTTCTTCAACGGTCCG 960
 QY 1285 ATTTGAGTGTGTACTGAGAGAGCTCCATTAAATTTGAGTAAAGCTGTTTCAACAAAGCG 1344
 Db 961 ATTTGAGTGTGTACTGAGAGAGCTCCATTAAATTTGAGTAAAGCTGTTTCAACAAAGCG 1020
 QY 1345 TCTTAACATTTGATGAAAGGTTTACCGCGCGCTATGTTCTCTGGAAGCTATTC 1404
 Db 1021 TCTTAACATTTGATGAAAGGTTTACCGCGCGCTATGTTCTCTGGAAGCTATTC 1080
 QY 1405 CAATGACATTCACCCAGAGTTTAAATTTAAATAAACCATTCGTTTCTGATGATCGAGC 1464
 Db 1081 CAATGACATTCACCCAGAGTTTAAATTTAAATAAACCATTCGTTTCTGATGATCGAGC 1140
 QY 1465 AGAACACTTAAAGCCCATTTGTTTATGGTAAAGTTTGTCAACCAACTCAGAGATGATCGA 1524
 Db 1141 AGAACACTTAAAGCCCATTTGTTTATGGTAAAGTTTGTCAACCAACTCAGAGATGATCGA 1200
 QY 1525 C 1525
 Db 1201 C 1201
 RESULT 7
 ABK88015
 ID ABK88015 standard; DNA; 1182 BP.
 XX
 AC ABK88015;
 XX
 DT 07-OCT-2002 (first entry)
 XX
 DE DNA encoding human alpha-1-antitrypsin (AAT) protein.
 XX Alpha-1-antitrypsin; AAT: human; gene; ds; protease inhibitor; malaria;
 KW emphysema; asthma; chronic obstructive pulmonary disease; eczema;
 KW cystic fibrosis; otitis media; otitis externa; HIV; psoriasis;
 KW human immunodeficiency virus; atopic dermatitis; muscular dystrophy;

[illegible]

QY 1476 AGCCATTGTTATGGTAAGTTGTCACACCACTCAGAACT 1518
 DB 1216 AGCCCCCTCTTCATGGGAGGTCGTCAACCCACGACAGAGT 1258

RESULT 9

AAQ89254
 ID AAQ89254 standard; cDNA; 1312 BP.

XX AAQ89254;
 XX 18-OCT-1995 (first entry)
 XX Human alpha-1-tryptsin cDNA.
 XX Alpha-1-tryptsin; protease-inhibitor; ss.
 XX Homo sapiens.

XX Key Location/Qualifiers
 FT CDS 28..1258
 FT sig_peptide /*tag= a
 FT mat_peptide /*tag= b
 FT /*tag= c

PN US5399684-A.

XX 21-MAR-1995.

PD 20-MAY-1982; 82US-0380310.

XX 20-MAY-1982; 82US-0380310.

PR 07-FEB-1984; 84US-0638980.

PR 03-MAR-1987; 87US-0022543.

PR 16-DEC-1987; 87US-0133190.

PR 16-SEP-1988; 88US-0246912.

PR 22-AUG-1989; 89US-0398288.

PR 11-MAR-1991; 91US-0666450.

PR 18-NOV-1992; 92US-0979556.

PR 02-JUL-1993; 93US-0086642.

XX (WASH-) WASHINGTON RES FOUND.

PI Davie EW, Kurachi K, Thirumalachary C, Woo SLC;

XX WPI; 1995-130740/17.

DR P-PSDB; AAR71969.

XX Human alpha-1-tryptsin (al-At) cDNA sequence - can be used for

XX the expression of al-At

XX Claim 1; Fig.1; 15pp; English.

XX The sequence of a human alpha-1-antitrypsin cDNA clone is given in

XX AAQ89254. Expression of the cDNA in host cell transformants has

XX allowed production of recombinant alpha-1-antitrypsin.

XX SQ Sequence 1312 BP; 339 A; 368 C; 324 G; 281 T; 0 other;

XX Query Match 28.6%; Score 436.4; DB 16; Length 1312;

XX Best Local Similarity 59.8%; Pred. No. 1.1e-99;

XX Matches 731; Conservative 0; Mismatches 491; Indels 0; Gaps 0;

QY 418 CAAATTTAGCCGAATTTGCTTTCTTTCTATAGACAATTAGCTCATCAAGTAATTCTTA 477
 DB 182 CCAACTTGGCTGAGTTGGCTTACGCCATATACGCCAGTGGCACACAGTCCACAGCA 241
 QY 478 CTAAACATTTTTTTAGTCTCTTCTATTGCCCACCTGCTTTGCCCATGTGTGATTTAGTGA 537
 DB 242 CCAATATCTTCTCTCCCACTGAGCATCGCTACAGCCCTTGCATGCTCTCCCTGGGGA 301
 QY 538 CTAAAGCCGATACCCATGACGAGATTTTGAAGGTTTAAACTTTAAATTTGACCGAATTC 597
 DB 302 CCAAGGCTCACACTCACGATGAATCTCTGGAGGCTGAATTTCAACCTCACGAGATTC 361
 QY 598 CAGAAGCCCAATTTACGAGAGGTTTTCAGAGAGTCTTTCAGAACTTTTGAATCAACTGAT 657
 DB 362 CGGAGGCTCAGATCCATGAGGCTTTCAGGAACCTCTCCGCTACCCCTCAACGAGCCAGCA 421
 QY 658 CTCATTTGCAATTAATACTACTGTTAAACGGTTTATTTTCTCTGAAGTTTAAATTTGGTGT 717
 DB 422 GCCAGCTCCAGCTGACCAACCGCAATGGCTTCTCTCAGCGAGGCTGAGCTAGTGG 481
 QY 718 ACAAAATTCCTAGAAGACGCTCAAGAACTATATCATAGTAGGCTTTTACCGTTAAATTTG 777
 DB 482 ATAAGTTTTGGAGGATGTTTAAAGTTGTACCACTCAGAAAGCTTCTCACTCTCAACTTCG 541
 QY 778 GTGATACTGAGGAAGCTAAAGCAAAATTAATGATTTATGTTGAGAAAGGACCCAGGTA 837
 DB 542 GGGACACCGAAGAGGCCAAGAACAGATCAACGATTACGTGGAGAGGGTACTCAAGGA 601
 QY 838 AGATCGTTGACCTAGTTAAAGAAATTAGATCGTATACCGTCTTGGCACTAGTTAACTATA 897
 DB 602 AAATTTGGATTTGGTCAAGGAGCTTGACAGAGACACAGTTTTTGTCTCTGCTGAATTACA 661
 QY 898 TTTTTTCAAGGTAAGTGGGAAGCTCTTTTCGAGGTTTAAAGTACTGAAGAGAGAGATT 957
 DB 662 TCTTCTTTAAAGGCAATGGGAGAGACCTTTTGAAGTCAAGGACACCGAGAGAGAGACT 721
 QY 958 TTCATGTTGATCAAGTTACTACTCTCAAAGTTCCTCAATGATGAAAGACTGGGTATGTTC 1017
 DB 722 TCCAGCTGGACCAAGTACCACCTGGAAGTGGCTGCTATGATGAAGCGTTTAGGCTAT 781
 QY 1018 ATATTCAACATTTGCAAAATTAAGTTCTTTGGTCTTTATTAATGAAGTATTTAGGTAACG 1077
 DB 782 ACATCCAGCATTTGAAGAGCTGTCCAGCTGGGTGCTGTGATGAATACCTGGGCAATG 841
 QY 1078 CTACTGCTATTTTTTTTTTACCAGACGAAGTAAAGCTTCAACATTTAGAGAAATGAGTTCA 1137
 DB 842 CCACCGCATCTCTCTCTGCTGATGAGGGGAACACTACAGCACTTGGAAATGAACTCA 901
 QY 1138 CTCATGACATTTACTTAATTTTATAGAACAGAGATCGTGTAGCGCTTCTCTGCACC 1197
 DB 902 CCCAGGATATCATCACCAGTTCTCTGGAAATGAAGACAGAAGGCTCTGCAGCTTACATT 961
 QY 1198 TGCCAAAGTTAAGTATATCACCAGTACTTACAGCTTAAATCTGTTTTAGGCCAGTTAGTGA 1257
 DB 962 TACCCAACTGTCATTTACTTGGAACTATGATCTGAAGAGCGTCTGTAGTCAACTGGGCA 1021
 QY 1258 TTACCAAGATTTTCTAAGGGTGGCGATTTGAGTGGTGTACTGAAGAGGCTCCATTAA 1317
 DB 1022 TCATAAGGTCTTTCAGCAATGGGGCTGACCTCTCCGGGGTACAGAGAGGAGGCCCTGTA 1081
 QY 1318 AATTGAGTAAAGCTGTTCAAAAGCGCTCTTAACATTTGATCAAAAGGTTACCGAGGCGG 1377
 DB 1082 AGCTCTCCAGGCGGTGATAGGCTGTCTGACCATCGACGAGAAAGGAGTGAAGCTG 1141
 QY 1378 CCGGCGTATGTTCTCGGAAGCTATTCCAAATGAGCATTTCCACGAGAAGTTAAATTTAATA 1437
 DB 1142 CTGGGCGCATGTTTTTAGAGGCCATACCCATGTCTATCGCCCCGAGGTCAAGTTCAACA 1201
 QY 1438 AACCAATTCGTTTTCTGATGATCGAGCAGAACACTAAAGGCCATTTGTTATGGGTAGG 1497
 DB 1202 AACCCCTTTGCTTTCTTAATGATTGAACAAATACCAAGTCTCCCTCTTTCATGGGAAAG 1261

QY 1498 TTGTCACCACTCAGAGTA 1519
 Db 1262 TGGTGAATCCCAAAATA 1283

RESULT 10

AAV28471
 ID AAV28471 standard; cDNA; 1312 BP.

XX AC AAV28471;

XX DT 21-AUG-1998 (first entry)

XX DE Nucleotide sequence of the alpha-1-antitrypsin.

XX KW Human alpha-1-antitrypsin; ATR-1; antibody; ATR-1 deficiency; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers

FT CDS 28..1257

FT /tag= a

FT /product= "alpha-1-antitrypsin"

XX US5736379-A.

XX PD 07-APR-1998.

XX PF 07-JUN-1995; 95US-0479545.

XX PR 20-MAY-1982; 82US-0380310.

XX PR 07-FEB-1984; 84US-0638980.

XX PR 03-MAR-1987; 87US-0022543.

XX PR 15-DEC-1987; 87US-0133190.

XX PR 16-SEP-1988; 88US-0246912.

XX PR 22-AUG-1989; 89US-0398288.

XX PR 11-MAR-1991; 91US-0666450.

XX PR 18-NOV-1992; 92US-0979556.

XX PR 02-JUL-1993; 93US-0086442.

XX PR 12-DEC-1994; 94US-0361689.

XX PA (WASH-) WASHINGTON RES FOUND.

XX PI Davie EW, Kurachi K, Thirumalachary C, Woo SLG;

XX DR WPI; 1998-239214/21.

XX DR P-PSDB; AAW56709.

XX PT DNA encoding alpha-1 anti-tryptsin - useful for, e.g. producing

XX PS recombiant alpha-1 anti-tryptsin

XX PS Claim 1; Fig 1; 15pp; English.

XX CC This is the nucleotide sequence encoding the novel human

XX CC alpha-1-antitrypsin (ATR-1) protein. Its products are useful for

XX CC producing recombinant ATR-1 polypeptides, which can be used to prepare

XX CC antibodies for detecting ATR-1 variants in the blood, as ligands in

XX CC assays for ATR-1, and to treat ATR-1 deficiency.

XX XX Sequence 1312 BP; 339 A; 368 G; 281 T; 0 other;

XX XX Best Match 28.6%; Score 436.4; DB 19; Length 1312;

XX XX Best Local Similarity 59.8%; Pred. No. 1.1e-99;

XX XX Matches 731; Conservative 0; Mismatches 491; Indels 0; Gaps 0;

QY 298 TGTGTGGTAACTCCCTGTTTCCAGTCAAGGCCATGGAAGACCCCTCAAGGCCGCG 357

Db 62 TGGCAGGCTGTGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCT 121

QY 358 CTCACAAAACACGACGACGACGACGACGACGACGACGACGACGACGACGACGAC 417

Db 122 CCAGAGACAGATACATCCCAACCATGATCAGGATCAGGATCAGGATCAGGATCAG 181

QY 418 CAAATTTAGCCGAATTTGCTTTTCTTTTGTATAGACAAATTAGCTCATCAAGTAATTTCTA 477
 Db 182 CCAACTTGGCTGAGTTGCGCTTACGCTTATACCGCAGCTGGCACACGAGTCCCAACAGCA 241
 QY 478 CTAACATTTTCTTTAGTCTGCTTTCTATTGCACTGCTTTTCCGCAATGTTGAGTTAGGTA 537
 Db 242 CCAATATCTTCTTCCCGCAGTGACATCGCTACAGCTTTGCAATGCTCTCCCTGGGA 301
 QY 538 CTAAGCCGATACCCATGACGAGATTTTGAAGGTTTAACTTTAACTTTGACCCAAATTC 597
 Db 302 CCAAGGCTCAGACTCAGATGAAATCTTGGAGGCTGAAATTTCAACCTCAGGAGATTC 361
 QY 598 CAGAAGCCCAATTTACGAGGCTTTTCAAGAGTTCTTGAAGAACTTTTGAATCAACTGATT 657
 Db 362 CGAGGCTCAGATCCATGAAGCTTCCAGAACTCTCCGTAACCTCAACGAGGAGTTC 421
 QY 658 CTCATTTGCAATTAACCTACTGTAACGCTTTTATTTTGTCTGAAGTTTAAATTTGGTTG 717
 Db 422 GCCAGCTCCAGTGCACCCGCAATGGCTGTCTCTCAGCGAGGCTGAAAGTCTAGTGG 481
 QY 718 ACAAATTCCTAGAGAGCTCAAGAACTATATCATAGTAGGAGCTTTTACCGTTAATTTTG 777
 Db 482 ATAAGTTTGGAGGATGTTAAAGTTTGTACCACTCAGAAAGCTTCACTGTCAACTTCG 541
 QY 778 GTGATCTAGAGAGCTAAAGAAATTAATGATTATTGTTGAGAAAGGACCCAGGTA 837
 Db 542 GGGACACCGAAGAGGCCAAGAAACAGATCAAGATTACGTGGAGAGGTTACTCAAGGA 601
 QY 838 AGATCGTTGACCTAGTTAAAGAAATTAGATCGTATACCGCTTTCCGACTAGTTAACTATA 897
 Db 602 AATTGTGGATTTGGTCAAGGAGCTTGACAGAGACACAGCTTTTGTCTGTGTAATACA 661
 QY 898 TTTTTCAGAGGTAAGTGGGACGCTTTCGAGGTTAAAGTAACTGTAAGAGGAGGATT 957
 Db 662 TCTTCTTTAAAGCAATGGGAGAGACCCCTTTGAAGTCAAGGACACCCGAGGAGGAGGACT 721
 QY 958 TTCAATGTTGATCAAGTTTACTTGTCAAAAGTTTCCAAATGATGAAAGACTGGGTATGCTCA 1017
 Db 722 TCCAGCTGGACAGGTCACCCGCTGAGGCTGCTATGATGAAGCTTTAGGCATGTTTA 781
 QY 1018 ATATTCAACATTCGAAAAATTAAGTTCTTGGTCTTTATTAAGAGTATTAGTTAAGG 1077
 Db 782 ACATCCAGCATTTGTAAGAAGCTGTCCAGCTGGGTGCTGTGATGAATACCTGGGCAATG 841
 QY 1078 CTACTGCTATTTTCTTTTACCGAGCAAGGTAAGCTTCAACATTTAGAGAATAGTTGA 1137
 Db 842 CCACGGCATCTTCTTCTGCTGATGAGGGGAACTACAGCACTGGAAATGAACTCA 901
 QY 1138 CTCATGACATTATTACTAAATTTTGAAGAACGAGGATCTCTAGGCTTCTCTGCACC 1197
 Db 902 CCCAGATATCATCCCAAGTTCTCTGGAAATGAAGACAGAGGCTCTGCCAGCTTACATT 961
 QY 1198 TGGCAAGTTAAGTATCACCGGTACTTACGACTTAAATCTGTTTGGCCAGCTTAGGTA 1257
 Db 962 TACCCAACTCTCCATTTACTTGAAGCTTATGATCTGAAGAGCGCTCTAGGCTCACTGGCA 1021
 QY 1258 TTACCAAGTTTCTTCAACGCTGCGATTTGAGTGGTGTACTGGAAGAGCTCCATTAA 1317
 Db 1022 TCATTAAGTCTTTCAGCAATGGGCTGACCTCTCCGGGTCACAGAGGAGGACCCCTGA 1081
 QY 1318 AATTGAGTAAAGCTGTTTCAAAAGCCGCTTTAACTATTGATGAAAGGGTACCAGGCCG 1377
 Db 1082 AGCTCTCAAGGCCGTGCTAAGGCTGTGCTGACCATCGAGGAGAAAGGAGCTGAAGCTG 1141
 QY 1378 CCGGGCTGATGTTCTTGAAGCTATTTCAATGAGCTATTTCCACCAAGAGTTTAAATTA 1437
 Db 1142 CTGGGGCCATGTTTTCAGAGCCATACCCATGCTATCCCGCCCGAGGTCAAGTTCAACA 1201
 QY 1438 AACCAATTCGTTTCTGATGATCGAGCAGAACTATAAAGCCCAATTTTATGGTGAAG 1497
 Db 1202 AACCTTTGCTCTTATTGATTGAACAAATACCAAGTCTCCCTCTCTCATGGGAAAG 1261
 QY 1498 TTGTCAACCCAACTCAGAGTA 1519

1262 TGGTGAATCCCAAAAATA 1283

RESULT 12

AAS45052
ID AAS45052 standard: CDNA: 1367 BP.

ID FMS45052
 XX
 AC AAS45052:

18-DEC-2001 (first entry)

XX cDNA encoding novel human secretory protein, Seq ID No 133.

XX Human; secreted protein; arthritis; Crohn's disease; sepsis; shock;
KW Ischaemia-reperfusion injury; haematopoiesis; cancer; neuropathy;
KW transgenic animal; Alzheimer's disease; Parkinson's disease; burn;
KW ankyrotrophic lateral sclerosis; platelet disorder; thrombocytopenia;
KW ulcer; osteoporosis; bone degenerative disorder; periodontal disease;
KW gut protection; lung; liver fibrosis; immune deficiency; infection;
KW severe combined immunodeficiency; SCID; autoimmune disorder; allergy
KW multiple sclerosis; rheumatoid arthritis; diabetes mellitus; asthma;
KW fertility; analgesic; pain; antigen; ss.

NS Homo sapiens.
XX
KW

XX
PN WO200166689-A2.

13-SEP-2001.

05-MAR-2001; 2001WO-US04942.

07-MAR-2000; 2000US-0519705.

PR 19-MAY-2000; 2000US-0574454.
PR 17-JUN-2000; 2000US-0596193.

14-JUL-2000; 2000US-0616847.
PR
19-SEP-2000; 2000US-0665363.
PR

PR 20-OCT-2000; 2000US-0693267. XX

PA (HYSE-) HYSEQ INC.
XX

PI	Tang YT,	Liu C,	Asundi V,
PI	Zhao QA,	Yang Y,	Drmanac R

XX
DR WPI; 2001-589934/6

DR P-PSDB; AAU28152.
XX
XX

PT	Novel polypeptide prepared from var
PT	
PT	

PT cancer, neurological, inflammatory, and autoimmune disorders

PS Claim 1; SEQ ID No 133; 107pp; English.
XX

The invention relates to novel isolated human secreted polypeptides (I), (II) and polynucleotides (III). (I) and (II) are useful for treating inflammatory conditions such as arthritis, nephritis, Crohn's disease, ischaemia-reperfusion injury, shock, sepsis, immune responses, and is involved in increasing haematopoiesis, stem cell survival, bone growth and remodeling. (I), (II) and modulators of (II) are useful for prophylaxis or treatment of one or more cancers. (II) is also useful for creating transgenic animals useful for studying the *in vivo* activities of the polypeptide as well as for studying modulators of the polypeptides. (I) induces the proliferation of neural cells and regeneration of nerve and brain tissue and is useful for the treatment of central and peripheral nervous system diseases and neuropathies, such as Alzheimer's, Parkinson's disease, Huntington's disease, and amyotrophic lateral sclerosis. In addition, (I) is involved in chemotactic or chemokinetic activity, regulation of haematopoiesis and is useful for treating myeloid or lymphoid cell disorders, platelet disorders such as thrombocytopenia and for regeneration of bone, cartilage, tendon, ligament and/or nerve tissue growth, and in tissue repair, healing of burns, incisions, ulcers, for treating osteoporosis, osteoarthritis, bone degenerative disorders, or periodontal disease. Furthermore, (I) is also useful for

Db 894 CCCACGATATCATCCCAAGTCTCTGAAAATGAAGACAGAAGGTCTGCCAGCTTACAT 953
QY 1198 TGCCAAAGTTAAGTATCACCAGTACTAGACTTAAATCTGTTTATAGCCAGTTAGGTA 1257
Db 954 TACCCAAACTGTCCATTAAGTCTGGAACCTATGATCTGAAGAGCGTCTGGGTCAACTGGGA 1013
QY 1258 TTACCAAGTTTCTTCTAAGGTCGCGATTGAGTGGTCTTACTGAAGAGCTCCCATTTAA 1317
Db 1014 TCACCTAAGGTCCTTACGAATGGGCTGAGCTCTCCGGGTACAGAGGAGGACCCCTGA 1073
QY 1318 AATGAGTAAAGCTGTTCACAAAGCCGTCTTAACTATGATGAAAAGGTACGGAGCCG 1377
Db 1074 AGCTCTCCAAAGCGGTGCATGAAGCTGTGCTGACCATCGACGAGAGAAAGGACTGAAGCTG 1133
QY 1378 CCGGCGTATGTTCTTCTGGAAGCTATCCCATGAGCATTCACACAGAAAGTTAAATTTAAATA 1437
Db 1134 CTGGGCGCATGTTTATAGAGGCCATACCCATGCTCTATCCCCCGAGGTCAAGTTCAACA 1193
QY 1438 AACCATTCGTTTCTGATGATGAGCAGACAACTAAAAGCCCATTTGTTATGGGTAAGG 1497
Db 1194 AACCTTTGTTCTTAAATGATTGAACAAAATACCAAGTCTCCCTCTTTCATGGGAAAAG 1253
QY 1498 TTGTCACCCCAACTCAGAAGTA 1519
Db 1254 TGGTGAATCCCAACCAAAAATA 1275

RESULT 14
ABL67511
ID ABL67511 standard; DNA; 1352 BP.
XX
AC ABL67511;
XX
DT 15-MAY-2002 (first entry)
XX
DE Thyroid cancer related gene sequence SEQ ID NO:5848.
XX
KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
KW cytostatic; gene therapy; antineoplastic; Wilms' tumour; adenocarcinoma;
KW gene; ds.
XX
OS Homo sapiens.
XX
PN WO200194629-A2.
XX
PD 13-DEC-2001.
XX
PF 30-MAY-2001; 2001WO-US10838.
XX
PR 05-JUN-2000; 2000US-209473P.
PR 05-JUN-2000; 2000US-209531P.
PR 18-SEP-2000; 2000US-233133P.
PR 18-SEP-2000; 2000US-233617P.
PR 20-SEP-2000; 2000US-234009P.
PR 20-SEP-2000; 2000US-234034P.
PR 20-SEP-2000; 2000US-234052P.
PR 20-SEP-2000; 2000US-234509P.
PR 22-SEP-2000; 2000US-234567P.
PR 25-SEP-2000; 2000US-234923P.
PR 25-SEP-2000; 2000US-234924P.
PR 25-SEP-2000; 2000US-235077P.
PR 25-SEP-2000; 2000US-235082P.
PR 25-SEP-2000; 2000US-235134P.
PR 25-SEP-2000; 2000US-235280P.
PR 26-SEP-2000; 2000US-235637P.
PR 26-SEP-2000; 2000US-235638P.
PR 27-SEP-2000; 2000US-235711P.
PR 27-SEP-2000; 2000US-235720P.
PR 27-SEP-2000; 2000US-235840P.
PR 27-SEP-2000; 2000US-235863P.
PR 28-SEP-2000; 2000US-236028P.
PR 28-SEP-2000; 2000US-236032P.

28-SEP-2000; 2000US-236033P.
28-SEP-2000; 2000US-236034P.
28-SEP-2000; 2000US-236109P.
28-SEP-2000; 2000US-236111P.
29-SEP-2000; 2000US-236842P.
29-SEP-2000; 2000US-236891P.
02-OCT-2000; 2000US-237172P.
02-OCT-2000; 2000US-237173P.
02-OCT-2000; 2000US-237278P.
02-OCT-2000; 2000US-237294P.
02-OCT-2000; 2000US-237295P.
03-OCT-2000; 2000US-237316P.
03-OCT-2000; 2000US-237425P.
03-OCT-2000; 2000US-237598P.
03-OCT-2000; 2000US-237604P.
03-OCT-2000; 2000US-237606P.
03-OCT-2000; 2000US-237608P.
01-NOV-2000; 2000US-244867P.
01-NOV-2000; 2000US-245084P.
(AVAL-) AVALON PHARM.
Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
Soppet DR, Weaver Z;
WPI; 2002-188264/24.
Screening for anti-neoplastic agent involves exposing cells to a
chemical agent to be tested for anti-neoplastic activity, and
determining a change in expression of a gene of a signature gene set -
Claim 1; SEQ ID 5848; 44pp; English.
The present invention describes a method (M1) for screening for an
anti-neoplastic agent. The method involves exposing cells to a chemical
agent to be tested for anti-neoplastic activity, determining a change in
expression of at least one gene (I) of a signature gene set, where (I)
comprises a sequence (S) selected from 8447 sequences (given in ABL61664
to ABL70110), or is at least 95% identical to (S), where a change in
expression is indicative of anti-neoplastic activity. (I) has cytostatic
activity and can be used in gene therapy. M1 can be used for screening
an anti-neoplastic agent, and can be used for producing a product which
is the data collected with respect to the anti-neoplastic agent as a
result of M1, and the data is sufficient to convey the chemical
structure and/or properties of the agent. M1 can be used in the
treatment of cancer such as colon, breast, stomach, lung, thyroid,
oesophageal, ovarian, kidney, prostate or pancreatic cancer,
adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,
infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine
carcinoma, papillary carcinoma and Wilms' tumour.
Query Match 28.4%; Score 433.2; DB 24; Length 1352;
Best Local Similarity 59.7%; Pred. No. 6.8e-99;
Matches 729; Conservative 0; Mismatches 493; Indels 0; Gaps 0;
QY 298 TGTGTGTAAGTCTGTGTTTCCCGACGTCATGGAAGACCCCTCAAGGGGAGCGCG 357
Db 54 TGGCAGGCGCTGTGCTGCCCTGTCTCCCTGGCTGAGGATCCCGAGGAGATGCTG 113
QY 358 CTCAAAAAACCGACACCGACGATCATCAGCACCAAGACCATCCGACTTTTAATAAATTAATCTC 417
Db 114 CCAGAGACAGATACATCCACCATGATCAGGATCATCCCAACCTTCAACAAGATCACCC 173
QY 418 CAAATTTAGCGCAATTTGCTTTTCTTTGTATAGACAAATAGCTCATCAAGTAATCTA 477
Db 174 CCAACCTGGCTGAGTTCGCTTACGCCCTATACCCCGAGCTGGCACACCGATCCACAGCA 233
QY 478 CTAACATTTTCTTTAGTCTCTTCTTCTATTCGCACTGCTTCGCCCATTTGAGTTTAGGTA 537
Db 234 CCAATATCTTCTTCTCCCGACGATGAGCATCGCTACAGCGCTTGTGCAATGCTCTCCCTGGGGA 293

